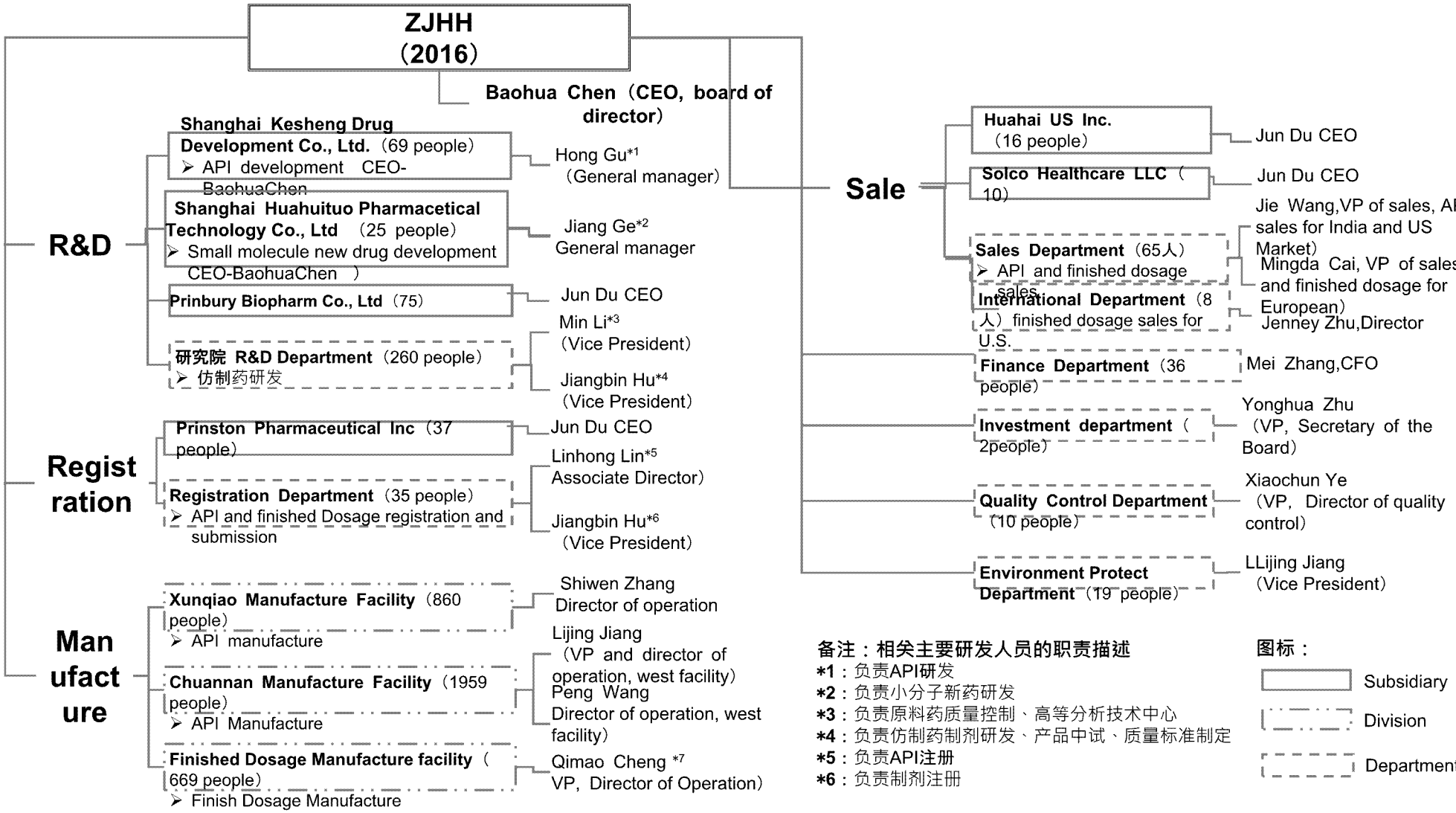
Exhibit B



备注：相关主要研发人员的职责描述

*1：负责API研发
*2：负责小分子新药研发
*3：负责原料药质量控制、高等分析技术中心
*4：负责仿制药制剂研发、产品中试、质量标准制定
*5：负责API注册
*6：负责制剂注册

图标：

Subsidiary
Division
Department

备注：R&D related description of role and responsibilities

***1**：负责API研发 – Responsible for API development

***2**：负责小分子新药研发 – Responsible for small molecule new drug development – Not related to US product

***3**：负责原料药质量控制、高等分析技术中心-Responsible for quality control of API, high analytical technology

***4**：负责仿制药制剂研发、产品中试、质量标准制定 – Responsible for the generic pharmaceutical preparations, product testing, setting quality standard –Not related to US Product

***5**：负责API注册 –Responsible for API registration

***6**：负责制剂注册 – Responsible for finished dosage-not related to US product

***7: Includes two R&D departments (Tech and quality control development) to work with US R&D to perform scale up and tech transfer analytical research and development**

Note: All the VP or directors are reported to Baohua Cheng Directly